



(For Professional Use Only) V6

1. Intended Purpose

The TANBead® Nucleic Acid Extraction Kit is a nucleic acid purification kit based on magnetic bead technology by using with corresponding TANBead® Nucleic Acid Extractor, which can automatically isolate and purify total RNA from a broad range of viruses, such as hepatitis C virus in human serum and samples suspended in phosphate buffered saline (PBS). The purified RNA can be used with any downstream application employing PCR-based qualitative, semi-quantitative and quantitative assays. The kit is intended for use by technicians, physicians, and biologists with well-trained in molecular biological techniques, the techniques of magnetic bead purification and in vitro diagnostic procedures. Any diagnostic results generated by using the sample preparation procedure in conjunction with any downstream diagnostic assay should be interpreted related to other clinical or laboratory findings. The kit is not limited to any specific disorder, condition, or other additional accompanying diagnostics. It is applicable for all population.

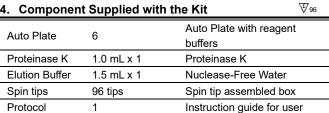
2. The basic principle

The silicon dioxide layer coated on the magnetic beads can adsorb the negatively charged molecules to purify nucleic acids from samples.

3. Specification

Starting Materials	300 μL serum or PBS		
	suspension		
Elution Volume	50 - 80 μL		

4. Component Supplied with the Kit



5. Auto Plate Content

Well	Buffer	Volume (μL)
1/7	Lysis Buffer	600
2/8	Washing Buffer 1	800
3/9	Washing Buffer 2	800
4 / 10	Washing Buffer 2	800
5 / 11	Magnetic Beads	800
6 / 12	Elution Buffer	80

6. Kit Storage and Shelf Life

- Components under room temperature (15~35°C) can be stored until the expiration date labeled on the box.
- 2) The proteinase K is transported at room temperature. Upon received, please store proteinase K at 2~8°C.

7. Precautions

- 1) It can only be used for in vitro diagnostic.
- 2) Avoid using expired reagents.
- 3) When the temperature is below 20°C, place the Auto Plates in an oven (preheated 42~60°C) 5 to 10 minutes.
- 4) Avoid vigorous shaking, in order to avoid excessive formation of
- 5) Carefully remove aluminum foil to avoid splashing.
- 6) Do not expose the opened reagents or Auto Plates to air. The evaporation would lead to pH change, or effect on the extraction effectiveness.
- 7) Please check the integrity of the Auto Plates and remember to mount the spin tips into the appropriate position of the suitable instrument before operating them.
- 8) Please wear a mask and disposable gloves when handling.
- Use sterile consumables to avoid nuclease contamination.
- 10) Reagent solution contains guanidine salt, avoid using bleach containing detergent.

- 11) Avoid eyes, skin, and clothing contact with reagents. In case of any contact, flush with flowing water.
- 12) If any serious incident occurs, please report to the manufacturer and the competent authority of the member state in which the user and / or the patient is established.
- 13) The reagents are all colorless and transparent. Colored reagents indicate contamination, please replace it with a fresh plate before proceeding.

8. Materials required, Not Supplied

- 1) TANBead® Nucleic Acid Extraction System Model: Maelstrom 8 series, Maelstrom 4800 series (non-sterile)
- 2) Disposable gloves
- 3) Scissors, utility knives
- 4) Micropipette, disposable tips (10 μ L / 200 μ L / 1000 μ L)
- 5) 1.5 mL microcentrifuge tube
- 6) 15 mL / 50 mL conical tube

9. Sample Collection, Transportation, and Storage

Sample collection and storage

- 1) Serum, whole blood
 - a. Serum specimens must be obtained from serum collection tubes, whole blood specimens must be obtained from sodium citrate or EDTA collection tubes.
 - b. Fresh whole blood specimens can be stored at room temperature for 6 hours.
 - c. After centrifugation, the serum sample can be stored at
 - i. Room temperature for 24 hours.
 - ii. 2~8°C up to 7 days.
 - iii. -20°C long-term preservation.

Specimen transportation

Transportation of whole blood, serum specimens should be followed by specific pathogen transportation-related laws. The whole blood sample should be kept between 2~25°C during transportation and within 6 hours for separated serum. Serum samples can be transported between 2~8°C or by freezing.

10. Nucleic Acids Extraction Protocol

- 1) Carefully remove the aluminum foil on the Auto Plates.
- 2) Add 300 µL serum or PBS suspension and 10 µL Proteinase K into well #1 / #7 of Auto Plate (Plate filled with lysis buffer). Note: The volume ratio of sample and lysis buffer is about 300 µL: 600 µL. Changing this ratio might affect the performance of this kit.
- Set up spin tips.

Maelstrom 8 series: Handle to mount tips and make sure that there is no gap between the necks of spin tips and the spin shaft. Maelstrom 4800 series: Go to Tip page and press the mount tips region.

- Push Auto Plates completely to the bottom of the plate rack. Make sure that the chamfer of the plate is at the lower left.
- Select the program

Maelstrom 8 series: Press "665-1" for input specimens at column #1 or "665-7" for input specimens at column #7.

Maelstrom 4800 series: "665" or "665-Rapid".

Note: Program "665-Rapid" improves the processing time. The actual extraction performance depends on various sample type.

The parameters are given in the following section.

- 6) Carefully remove the Auto Plates when the program is finished.
- 7) Use micropipette to transfer the purified nucleic acids from well #6 / #12 to a clean tube.
- 8) Discard used Auto Plates and spin tips into the waste recycling

11. Program

■ Maelstrom 8 series

Program Name: 665-1 / 7						
Well	1/7	2/8	3/9	4 / 10	5 / 11	6 / 12
Volume	900 (μL)	800 (μL)	800 (μL)	800 (μL)	800 (μL)	100 (μL)

Step	Well	Action	RPM	Time (Second)	CW/CCW (Second)	Temp.	Temp. Control
1	5/11	Collection	0	30	0	60	Yes
2	1/7	Mixing	3000	480	0	60	Yes
3	1/7	Collection	0	30	0	60	Yes
4	2/8	Mixing	3000	60	0	45	Yes
5	2/8	Collection	0	30	0	45	Yes
6	3/9	Mixing	3000	60	0	45	Yes
7	3/9	Collection	0	30	0	45	Yes
8	4 / 10	Mixing	3000	60	0	45	Yes
9	4/10	Collection	0	30	0	45	Yes
10	4 / 10	Vapor	0	300	0	45	Yes
11	6 / 12	Mixing	3000	300	0	45	Yes
12	6 / 12	Collection	0	30	0	45	Yes
13	5/11	Mixing	3000	60	0	0	No

■ Maelstrom 4800 series

Program N	lame: 665			Model: N	1aelstrom 4	800 series	
Temp1	Temp2						
40	40						
Well	Name	Volume	Action	Mixing	Collect		
1	LB	900	Rev. U / D	Low	Low		
2	WB1	800	For.	Low	Low		
3	WB2	800	For.	Low	Low		
4	WB2	800	For.	Low	Low		
5	MB	800	For.	Low	Low		
6	EB	150	For.	Low	Low		
Step	Well	Temp (°C)	Mixing (M)	Mixing Speed (RPM)	Collect (M)	Vapor (M)	Pause
1	5	-	0	2500	0.5	0	Off
2	1	55	12	1500	0.5	0	Off
3	2	-	2	2500	0.5	0	Off
4	3	-	1	1500	0.5	0	Off
5	4	-	1	1500	0.5	10	Off
6	6	Off	5	2500	1.5	0	Off
7	3	-	0.5	2500	0	0	Off

Program N	lame: 665-R	apid		Model: N	1aelstrom 4	800 series	
Temp1	Temp2						
Off	Off						
Well	Name	Volume	Action	Mixing	Collect		
1	LB	900	Rev. U/D	Low	Low		
2	WB1	800	For.	Low	Low		
3	WB2	800	For.	Low	Low		
4	WB2	800	For.	Low	Low		
5	MB	800	For.	Low	Low		
6	EB	150	For.	Low	Low		
Step	Well	Temp (°C)	Mixing (M)	Mixing Speed (RPM)	Collect (M)	Vapor (M)	Pause
1	5	-	0	3000	0.5	0	Off
2	1	60	8	3000	0.5	0	Off
3	2	-	1	3000	0.5	0	Off

Step	Well	Temp (°C)	Mixing (M)	Mixing Speed (RPM)	Collect (M)	Vapor (M)	Pause
4	3	-	1	3000	0.5	0	Off
5	4	-	1	3000	0.5	5	Off
6	6	45	5	3000	0.5	0	Off
7	3	-	0.2	3000	0	0	Off

12. Result

Nucleic acid product purified by TANBead® nucleic acid extraction kit can perform qualitative / quantitative analysis of specific genes by PCR, RT-PCR, Q-PCR or qRT-PCR. Please refer to the molecular diagnostic kit manual.

13. Reagent performance

■ Repeatability

Under repeatability conditions where nucleic acids are extracted with the same reagent kit on the same HCV serum concentration by the same operator. The coefficient of variation of nucleic acid extraction concentration is less than 5%.

■ Reproducibility

A five-day reproducibility test was carried out with 100 IU / mL of HCV serum samples for 5 consecutive days with the same reagent kit by different operators. The coefficient of variation of nucleic acid extraction concentration is less than 5%.

■ Detection limit of HCV virus: ≥100 IU / mL

■ Interfering substance

According to preclinical tests, the performance of extraction kit will not be affected by EDTA, Li-Heparin, Sodium Citrate, D-Glucose, Hemoglobin, lipoprotein and triglyceride in samples.

■ The stability of extracted RNA

Storage Conditions	RNA stability
-80°C	Over 90 days
-20°C	28 days
4°C	14 days
25°C	2 days
Freeze-thaw	10 times

14. Explanation of Symbols

***	Manufacturer	(i	Consult instructions for use
15°C- 35°C	Temperature limit	Σ	Contains sufficient for test
CE	CE mark	IVD	In vitro diagnostic medical use
REF	Catalogue number	\triangle	Caution
LOT	Batch code	NON	Non-sterile
(2)	Do not re-use	漆	Keep away from sunlight
س	Date of manufacture	8	Use-by date

EC REP

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15. Post-market surveillance conclusion

After a risk assessment and clinical evaluation assessment, when weighing the benefits of medical device, patients, and the risks associated with the use of the device, the risk is acceptable. The postmarket surveillance report shows that no death or serious adverse events occurred.

